

- 48. (New) A method of extracting glucosinolates and isothiocyanates from plant tissue comprising homogenizing said plant tissue in an excess of a mixture of dimethyl sulfoxide, acetonitrile and dimethylformamide at a temperature sufficient to inactivate myrosinase enzyme activity.
- 49. (New) The method of claim 48, wherein the ratio of dimethyl sulfoxide:acetonitrile:dimethylformamide is 1:1:1.
- 50. (New) The method of claim 48, wherein said temperature is between 0°C and the freezing temperature of the extraction mixture.
- 51. (New) The method of claim 48, wherein said temperature is between -50°C and the freezing temperature of the extraction mixture.
 - 52. (New) The method of claim 48, wherein said plant tissue is rich in glucosinolates.
- 53. (New) The method of claim 52, wherein said plant tissue is selected from the group consisting of cruciferous sprouts measured after 3 days of growth, cruciferous seeds, plants or plant parts.
- 54. (New) The method of claim 53, wherein said sprouts, seeds, plants or plant parts have at least 200,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 55. (New) The method of claim 53, wherein said sprouts, seeds, plants or plant parts have at least 300,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 56. (New) The method of claim 53, wherein said sprouts, seeds, plants or plant parts have at least 400,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 57. (New) The method of claim 53, wherein said sprouts, seeds, plants or plant parts have at least 500,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 58. (New) A method of extracting glucosinolates and isothiocyanates from plant tissue rich in glucosinolates, with the exception of cabbage, cress, mustard and radish sprouts, comprising homogenizing said plant tissue in a non-toxic solvent at a temperature sufficient to inactivate myrosinase enzyme activity.



ļ.

Cub cont.

- 59. (New) The method according to claim 58, wherein said solvent is water.
- 60. (New) The method of claim 59, wherein said water is 100°C.
- 61. (New) The method according to claim 58, wherein said solvent is liquid carbon dioxide.
 - 62. (New) The method according to claim 58, wherein said solvent is ethanol.
- 63. (New) The method of claim 58, wherein said plant tissue is selected from the group consisting of cruciferous sprouts measured after 3 days of growth, cruciferous seeds, plants and plant parts.
- 64. (New) The method of claim 63, wherein said sprouts, seeds, plants or plant parts have at least 200,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 65. (New) The method of claim 63, wherein said sprouts, seeds, plants or plant parts have at least 300,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 66. (New) The method of claim 63, wherein said sprouts, seeds, plants or plant parts have at least 400,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.
- 67. (New) The method of claim 63, wherein said sprouts, seeds, plants or plant parts have at least 500,000 units per gram fresh weight of Phase 2 enzyme-inducing potential.